Case report: a subject with a mutation in the ATG start codon of L-ferritin has no haematological or neurological symptoms

L Cremonesi, A Cozzi, D Girelli, F Ferrari, I Fermo, B Foglieni, S Levi, C Bozzini, M Camparini, M Ferrari, P Arosio

RESULTS AND DISCUSSION

DNA variations in the 5’UTR of L-ferritin were analysed in a cohort of 292 subjects who participated in a control study on the incidence of HHCS in subjects with cataracts. The participants consisted of 177 subjects with cataracts and 115 healthy controls. One of the subjects showed an abnormal DHPLC pattern (fig 1A) indicating the presence of a mutation. This was confirmed by sequence analysis that showed an A→G substitution causing the first Met codon to change into a Val (fig 1B). The L-ferritin transcript does not have any ATG upstream of the start codon and the improbable use of the next in-frame ATG (Met-69) would encode a non-functional and unstable protein of 106 amino acids. Therefore the A→G mutation was predicted to disable protein translation and expression. The mutation seems to be rare and was not found in an additional 1006 patients at risk for haemochromatosis or HHCS and screened for the same DNA region (not shown). The subject was a 52 year old female with keratoconus in the left eye. Her blood ferritin level was 1320 μg/l. She showed a 1A G mutation which alters the ATG start codon in a healthy control subject with normal haematological indices and normal haematological values. The mutation is predicted to disable protein translation and expression.

Key points

- Mutations of the L-ferritin gene (FTL) are associated with two types of dominant genetic disorder: hereditary hyperferritinaemia cataract syndrome caused by modifications of the iron responsive element in the 5’UTR which upregulate ferritin expression, and hereditary neuroferritinopathy caused by an adenine insertion at position 460–461 of the mRNA coding region.
- We report the identification of the heterozygous mutation 1A→G which alters the ATG start codon in a healthy control subject with normal haematological indices and no signs of neurological or movement disorders, but with low levels of L-ferritin in serum and in blood cells.
- The data indicate that haploinsufficiency of L-ferritin protein has no evident clinical effects on systemic iron metabolism, and suggest that the defects caused by the FTL mutation in neuroferritinopathy probably originate from abnormal properties of the altered protein.

Abbreviations: HHCS, hyperferritinaemia cataract syndrome; IRE, iron responsive element; IRP, iron regulatory protein.

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woman with no history of iron-deficient anaemia, pregnancy, regular blood donations, or gastrointestinal diseases associated with blood loss. Family history did not reveal any genetic disease, particularly with respect to neurological or movement disorders. The woman was in good health with the exception of mild hypertension treated with a low-dose β-blocker and diuretics. Haematological examinations showed normal levels of haemoglobin (14.4 g/dl), MCV (88.5 fl), and iron indices (serum iron 70 μg/l; transferrin saturation 17%), with the relevant exception of serum ferritin levels, which were low (12.7 μg/l). In agreement with normal haemoglobin and MCV, the normal levels of serum transferrin receptor (1.61 mg/l; range 0.8–1.8) ruled out the presence of true iron deficiency. A detailed family study could not be performed. To evaluate whether the low serum ferritin levels unrelated to body iron stores were caused by the genetic effect we measured ferritin levels in the most easily obtainable tissue, that is blood cells. In two independent determinations we found that H-ferritin concentrations in the subject and a healthy control were comparable (mean values 1000 v 1400 ng/g protein, respectively), while that of L-ferritin was greatly reduced in the subject (47 v 340 ng/g, respectively). The ratio between L- and H-ferritin in the control was 0.24, consistent with the figures reported by Piperno et al,16 while the index in the subject was 0.047, which is about five fold lower. For ethical reasons, brain MRI for investigation of iron deposition in basal ganglia could not be performed.

Present data indicate that the inactivation of one FTL allele causes a significant reduction of L-ferritin in serum and blood cells. This was expected, since mice with an inactivated H-ferritin allele showed low H-ferritin levels in all tested tissues.17 More important is the observation that genetic L-ferritin deficiency (in the present case) or excess (in HHCS) has no evident effect on systemic iron metabolism7 indicating that L-ferritin levels do not regulate iron homeostasis. In neuroferritinopathy all subjects with the L-ferritin mutation develop movement disorders typically in the fourth to fifth decades, with some as early as in the late teens, and the disease is considered fully penetrant by the age of 60.11–13 The 52 year old woman we described had no neurological or movement abnormalities and none were reported in her family. The observation suggests that the A→G substitution in codon 1 of L-ferritin did not cause the abnormal iron deposition in the basal ganglia of the brain typical of neuroferritinopathy, although this cannot be completely excluded as abnormal iron deposition might develop at a later age. This mutation is predicted to simply disable protein expression, while the A insertion found in neuroferritinopathy is predicted to cause the expression of a protein structurally different from wild type L-ferritin, which may interfere with ferritin functionality and with cellular iron homeostasis. In conclusion, the present case report suggests that neuroferritinopathy is not a consequence of haploinsufficiency of L-ferritin protein, but more likely a gain-of-function caused by the predicted abnormal C-terminus of the protein.

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